ST AVAILABLE COPY

PATENT SPECIFICATION

NO DRAWINGS



875,133

50

Date of Application and filing Complete Specification Nov. 14, 1957. No. 35552/57.

Application made in Italy on Nov. 15, 1956.

Complete Specification Published Aug. 16, 1961.

Index at acceptance: —Classes 2(6), P7(A:C10:D1X:S2); 2(2), B2B2.

International Classification: - C08f. D01f.

COMPLETE SPECIFICATION

Preparation of Uniformly Dyed Elongated Articles of Polypropylene

We, Montecatini Societa Generale Per L'Industria Mineraria E Chimica, a Body Corporate organised and existing under the laws of Italy, of 18, Via Filippo Turati, Milan, Italy, do hereby declare this invention, for which we pray that a patent may be granted to us, and the method by which it is to be performed, to be particularly described in and by the following statement:—

The present invention relates to the preparation of uniformly dyed or coloured elongated articles formed predominantly from highly crystalline polypropylene, that is to say articles such as yarns, fibres, tapes and films formed predominantly of highly crystalline polypropylene which have one or both of their transverse dimensions small compared with their length dimensions. Such articles may be produced for example by extrusion or spinning.

It is known that highly crystalline polypropylene is a colourless material with a high melting point and insoluble in the common solvents, which although it can be formed into elongated articles having very useful physical, chemical and mechanical properties, has a very low affinity for dyes.

We have now found that, in the case of polypropylene, it is possible to obtain uniformly dyed manufactured articles by having recourse to a bulk dyeing process.

The invention provides a process for the preparation of uniformly dyed elongated articles formed predominantly of highly crystalline polypropylene which process comprises mixing at least one dye which is stable at the melting point of the polypropylene and does not effect a degrading action thereon, with the polypropylene, melting or dissolving the polypropylene and subsequently forming the elongated articles.

The article may be formed by an extrusion

or spinning process and either organic dyes or inorganic pigments may be used.

The dye may be mixed with polypropylene 45 in any of several ways:—

1) It can be introduced before the poly-

propylene is melted, by mixing it intimately with polypropylene in powdered form.

2) It can be introduced into the polypropy-

lene after the polypropylene has been melted or is in the form of a solution or

3) The dve and relevant length of the solution or

 The dye and polypropylene may be dissolved in a common solvent.

4) The dye and polypropylene may be dissolved in a mixture of two or more solvents at least one of which is a solvent for the dye.

Suitable organic dyes are anthraquinone, aminoanthraquinone, indigo, thioindigo, phthalocyanine and azo dyes. In general, both organic dyes and inorganic pigments used must have the following essential properties, namely, stability at the melting point of polypropylene and a non-degrading action towards the polypropylene.

Mixtures of various dyes may be used to obtain particular dyeing effects provided that each component has these properties.

The dyed articles prepared by the process of this invention show a uniform distribution of the dye (as can be seen from examining thin sections under the microscope), and colours that are fast to light, washing and rubbing. Moreover, the dyes have no or only a very low tendency to migrate towards the polypropylene surface. Such a tendency which is common with many polymers tends considerably to reduce abrasion resistance.

The following examples are given to illustrate the invention:—

Example 1

5 g of a powdered dye of the anthraquinone series (Romanthrene yellow GCN) are added to 500 g of finely powdered crystalline poly-

[Price 3s. 6d.]

Price 2

	propylene. The mixture is introduced into a Werner type mixer and mixed for 30 minutes. The mixture is melted at 250° C. and then	shows the following characteristics:— tenacity 4.7 g/den	20
5	extruded through a spinneret having 18 holes of 0.3 mm diameter. The filaments thus obtained are stretched	The colour-fastness tests, carried out according to the specifications of the "International	
10		(dye: Romanthrene yellow GCN; extrusion temperature: 230° C; permanence time: 7 minutes)	
	tests and gives the following results:— tenacity 4.6 g/den elongation 25%. These data compared with those determined	natural light 6 washing 5 perspiration 5 fulling 5	30
15	on a filament prepared from a polymer having	trichlorethylene 5	
	the same physical characteristics of the test-		
	polymer but without any dye, show that the dye causes no degradation of polypropy-	in the same way employing the following dyes:	23
	Indanthren Scarler 2G	(anthraquinone series)	
	Durindone Scarlet YP Polymon blue GS	(thioindigo series) (phthalocyanine series)	
40	The colours obtained have good intensity and uniformity and the serimetrical	characteristics are the following:-	
	Indanthren Scarlet 2G	tenacity 4.92 g/den	
45	Durindone Scarlet YP	elongation 23% tenacity 4.9 g/den	
	Pol-man blue CS	elongation 22.5%	
	Polymon blue GS	tenacity 4.89 g/den elongation 23.5%	
	The colour-fasmess is as follows:—	The following dyes were also used:— Indanthrene Scarlet 2G	•
50	Indanthren Scarlet 2G	Durindone Scarler YP	80
	fastness to natural light 5	Polymon blue GS.	
	fastness to washing 5 fastness to perspiration 5 fastness to rubbing 5 fastness to fulling 5	The extrusion temperature is 250° C. Film	
	fastness to perspiration 5 fastness to rubbing 5	sections are cut from each, which, under the microscope, show no portions of clotted dye	
55	fastness to fulling 5	and are very transparent.	85
	fastness to trichlorethylene 5	Example 3	
	Demonson County ND	A solution is prepared by dispersing in the	
	DURINDONE SCARLET YP fastness to natural light 5	cold 18 parts by weight of crystalline poly- propylene having an intrinsic viscosity of 2.6	
	fastness to washing 5	(determined in a 1% by weight solution of	90
60	fastness to perspiration 5	tetrahydronaphthalene at 135° C.), in 62 parts	30
	fastness to rubbing 4	by weight of a petroleum fraction boiling at	
	fastness to fulling 5	180—200° C. 20 parts by weight cyclo-	
	fastness to trichlorethylene 5	hexanone containing 0.36 parts by weight	~=
	POLYMON BLUE GS	dissolved Oracet B blue are added. The whole is heated to 150° C. while stirring. The mix-	95
65		ture is then filtered through a filter-press and	
0,5	fastness to washing 5	extruded through a spinneret having 30 holes	
	fastness to perspiration 5	of 200μ diameter, into an evaporation tower	
	fasmess to rubbing 5	with circulating air which removes the sol-	100
70	fastness to natural light 5 fastness to washing 5 fastness to perspiration 5 fastness to rubbing 5 fastness to fulling 5 fastness to trichlorethylene 5	vent vapours. Dowtherm (Registered Trade	
70	fastness to trichlorethylene 5	Mark) at 300° is circulated in the tower jacket. The solution is delivered from the spin-	
	Example 2	neret in the form of filaments at a rate of	
	A mixture of 500 g powdered crystalline	25 m/minute and after a 6 m run in the	105
	polypropylene and 5 g Romanthrene GCN	evaporation tower, is wound up on a hobbin	
	yellow is intimately mixed in a Werner type	at a rate of 125 m/min. The varn obtained	
75	mixer for 30 minutes. The mixture is melted	is stretched with a 1:5 ratio at 145° C. in	
	and extruded in order to obtain a uniformly dyed film.	warm air. The yarn has the following characteristics:—	110
	UYCU IIIII.	LVA AUGUVU .	110

	tenacity 5.4 g/den elongation 22%	lene, melting or dissolving the polypropylene and subsequently forming the elongated	
5	intensive and uniform colour under the microscope. Colour fastness:— to natural light 6/7 to washing 5 to rubbing 5	articles. 2. A process according to Claim 1, wherein the article is formed by extrusion. 3. A process according to Claim 1, wherein the article is formed by spinning.	45
10	to perspiration 5 The intrinsic viscosity, determined as pre- viously described on the yarn dissolved in tetrahydronaphthalene, is 2.54.	4. A process according to any one of Claims 1 to 3, wherein the dye is an organic dye. 5. A process according to any one of Claims 1 to 3, wherein the dye is an inorganic pigment.	50
15	with one or more of the processes described in several of our copending applications. For	6. A process according to any one of the preceding claims, wherein the dye is introduced into the polypropylene in powdered form before the polypropylene is melted. 7. A process according to any one of	5 5
20	example stabilizing treatment as described in Application No. 22342/57 (Serial No. 813,891), filtering procedure as described in Application No. 27758/57 (Serial No. 827,424), cooling procedure as described in	into the polypropylene after the polypropylene has been melted. 8. A process according to any one of Claims	60
25	Application No. 31404/57 (Serial No. 875,132), the procedure of stretching in the presence of a plasticiser or swelling agent as described in Application No. 33069/57 (Serial No. 817,125), the spreading and	1 to 5, wherein the dye and polypropylene are dissolved in a common solvent. 9. A process according to any one of Claims 1 to 5 in which the dye and the polypropylene are dissolved in a mixture of two or more solvents at least one of which is a sol-	65
30	volarilisation procedure as described in Application No. 9676/57 (Serial No. 853,637), or the thermal degradation procedure described in Application No. 36349/56 (Serial No. 835,038).	vent for the dye. 10. A process for the preparation of uniformly dyed elongated articles substantially as herein described and illustrated by the fore-	70
35	WHAT WE CLAIM IS:— 1. A process for the preparation of uniformly dyed elongated articles formed predominantly of highly crystalline polypropy-	going examples. 11. Uniformly dyed elongated articles when prepared by the process claimed in any one of the preceding claims.	75
40	lene which process comprises mixing at least one dye which is stable at the melting point of the polypropylene and does not effect a degrading action thereon, with the polypropy-	ERIC POTTER AND CLARKSON, Chartered Patent Agents, 317, High Holborn, London, W.C.1.	

Learnington Spa: Printed for Her Majesty's Stationery Office, by the Courier Press.—1961.
Published at The Patent Office, 25, Southampton Buildings, London, W.C.2, from which copies may be obtained.